

4132 FIRST AID KIT - 4132 first aid kit

Honeywell Safety Products USA, INC

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, [click here](#).

4132 First Aid Kit (Eye Wash, BCM:FABC, BZK, Aypnal, sting rel, neomycin- SF00003655)

Active ingredient

Sterile Water 99%

Purpose

Eyewash

Uses

- For flushing the eye to remove loose foreign material, air pollutants, or chlorinated water

Warnings

For external use only- Obtain immediate medical treatment for all open wounds in or near the eyes. To avoid contamination, do not touch tip of container to any surface. Do not reuse. Once opened, discard.

Do not use

- if solution changes color or becomes cloudy
- if you have open wounds in or near the eyes, get medical help right away

Stop use and ask a doctor if

- you experience eye pain
- changes in vision
- continued redness or irritation of the eye
- condition worsens or persists

Keep out of reach of children

If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- remove contacts before using
- twist top to remove
- flush the affected area as needed
- control rate of flow by pressure on the bottle
- if necessary, continue flushing with emergency eyewash or shower

Inactive Ingredients

sodium chloride, sodium phosphate dibasic, sodium phosphate monobasic

Questions?

Call **1-800-430-5490**

Honeywell Safety Products USA, Inc. Smithfield, RI 02917

First Aid Burn Cream

Active ingredient

Benzalkonium chloride 0.13%

Lidocaine HCl 0.5%

First Aid Burn Cream

Purpose

First aid antiseptic

External analgesic

First Aid Burn Cream

Uses

- prevent skin infection
- for temporary relief of pain associated with minor burns

First Aid Burn Cream

Warnings

For external use only

Do not use

- in or near the eyes
- if you are allergic to any of the ingredients
- in large areas of the body, particularly over raw surfaces or blistered areas
- for more than 10 days

Ask a doctor before use if you have

- deep or puncture wounds
- animal bites
- serious burns

Stop use and ask a doctor if

- condition worsens
- symptoms persist for more than 7 days or clear up and occurs again within a few days

First Aid Burn Cream

Directions

- **adults and children 2 years of age and older:**
- clean the affected area
- apply a small amount of this product (equal to the surface area of the tip of a finger) onto affected area 1 to 3 times daily
- may be covered with a sterile bandage
- children under 2 years of age: consult a doctor

Other information

- tamper evident sealed packets
- do not use if packet is opened or torn

Inactive ingredients

aloe barbadensis juice, cetyl alcohol, diazolidinyl urea, edetate disodium, glycerin, glyceryl stearate SE, methylparaben, mineral oil, PEG-100, propylene glycol, propylparaben, stearic acid, trolamine, water

Questions

1-800-430-5490

BZK Antiseptic Wipe***Active ingredient***

Benzalkonium chloride 0.13%

BZK***Purpose***

First aid antiseptic

BZK***Uses***

Antiseptic cleansing of face, hands, and body without soap and water

BZK***Warnings***

For external use only

BZK***Do not use***

- in the eyes or over large areas of the body
- on mucous membranes
- on irritated skin
- in case of deep puncture wounds, animal bites or serious burns, consult a doctor
- longer than 1 week unless directed by a doctor

Stop use and ask a doctor if

- if irritation, redness or other symptoms develop
- the condition persists or gets worse

Keep out of reach of children

If swallowed, get medical help or contact a Poison Control Center right away.

BZK***Directions***

- .tear open packet and use as a washcloth

BZK

Other information

- store at room temperature 15 ° to 30 ° C (59 ° - 86 °F)
- do not reuse towelette

BZK

Inactive ingredients

water

BZK

Questions

1-800-430-5490

Aypanal

Active ingredient

Acetaminophen 325 mg

Aypanal

Purpose

Pain reliever/fever reducer

Aypanal

Uses

- temporarily relieves minor aches and pains due to the common cold and headache
- temporarily reduces fever

Ask a doctor before use if you have

liver disease

Aypanal

Warnings

Liver warning: This product contains acetaminophen. Severe liver damage may occur if you take: more than 4,000 mg in 24 hours, which is the maximum daily amount - child takes

Allergy alert: Acetaminophen may cause severe skin reactions. Symptoms may include:

skin reddening
blisters
rash

Do not use

- with any other drug containing acetaminophen (prescription or nonprescription). If you are not sure whether a drug contains acetaminophen, ask a doctor or pharmacist.

Ask a doctor before use if you have

liver disease

Ask a doctor or pharmacist before use if

you are taking the blood thinning drug warfarin

if pregnant or breast feeding

ask a health professional before use

Keep out of reach of children

Keep out of reach of children

Overdose Warning

Overdose warning: In case of accidental overdose, get medical help or contact a Poison Control Center right away. Prompt medical attention is critical for adults as well as for children even if you do not notice any signs or symptoms.

Aypanal

Directions

- do not take more than directed (see overdose warning) adults and children 12 years of age or older
- take two tablets every 4-6 hours while symptoms last
- do not take more than directed (see overdose warning)

adults and children 12 years of age or older

- take two tablets every 4-6 hours while symptoms last
- do not take more than 12 tablets in 24 hours
- children 6 to under 12 years of age
- take 1 tablet every 4-6 hours while symptoms last
- do not take more than 5 tablets in 24 hours
- children under 6 years
- consult a doctor

Aypanal

Other information

store at room temperature 15 ° to 30 ° C (59 ° - 86 °F) TAMPER EVIDENT PACKETS- DO NOT USE IF OPEN OR TORN

Aypanal

Inactive ingredients

corn starch, microcrystalline cellulose, povidone, sodium starch glycolate, stearic acid.

Aypanal

Questions or Comments?

1-800-430-5490

Sting Relief

Active ingredient

Ethyl alcohol 50.0%

Lidocaine HCl 2.0%

Sting Relief

Purpose

Antiseptic

Topical pain relief

Sting Relief

Uses

- prevent infection in minor scrapes, and temporary relief of itching of insect bites

Sting Relief

Warnings

For external use only

Flammable, keep away from open fire or flame

Do not use

- over large areas of the body
- in eyes
- over raw or blistered areas

Stop use and ask a doctor

- if conditions worsen or persist for more than 7 days or clear up and occur again within a few days

Keep out of reach of children

If swallowed, get medical help or contact a Poison Control Center right away

Sting Relief

Directions

- **adults and children 2 years and older:** Apply to cleaned affected area not more than 3 times daily.
- children under 2 years of age: consult a doctor.

Sting Relief

Inactive ingredients

benzalkonium chloride, menthol, and purified water

Sting relief

Questions or Comments

1-800-430-5490

Neomycin Antibiotic Ointment

Active ingredient

Neomycin sulfate (5 mg equivalent to 3.5 mg Neomycin base)

Neomycin Antibiotic Ointment

Purpose

First aid antibiotic

Neomycin Antibiotic Ointment

Uses

first aid to help prevent infection in - minor cuts - scrapes - burns

Neomycin Antibiotic Ointment

Warnings

For external use only

Do not use

- in the eyes
- over large areas of the body

Ask a doctor before use if you have

- deep or puncture wounds
- animal bites
- serious burns

Stop use and ask a doctor if

- the condition persists or gets worse
- a rash or other allergic reaction develops
- you need to use longer than 1 week

Keep out of reach of children

If swallowed, get medical help or contact a Poison Control Center right away

Neomycin Antibiotic Ointment

Directions

- clean the affected area
- apply a small amount of the product (an amount equal to the surface area of the tip of a finger) on the area 1 to 3 times daily
- may be covered with a sterile bandage

Neomycin Antibiotic Ointment

Other information

store at 15 ° to 25 °C (59 ° to 77 °F)

Neomycin Antibiotic Ointment

Inactive ingredient

petrolatum

Neomycin Antibiotic Ointment

Questions

1-800-430-5490

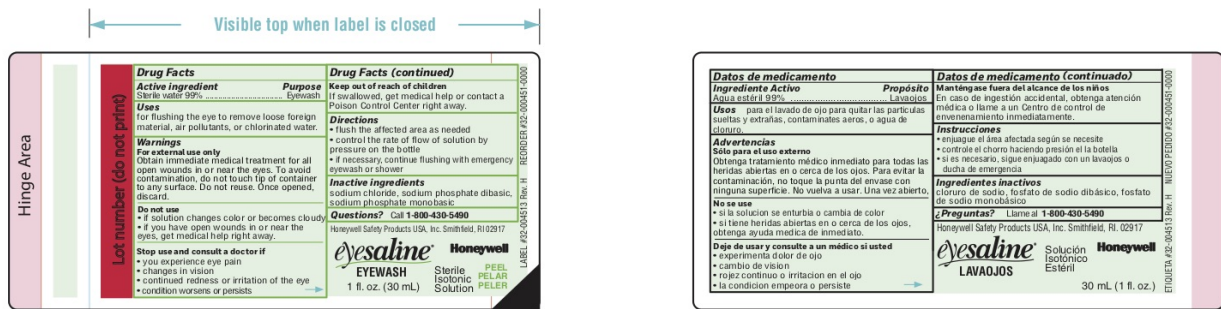
4132

SF00003655 Kit Contents

1 GAUZE PADS, 3" X 3", 4 PER
1 TWEEZER PLASTICS 4"
1 FIRST AID GUIDE ASHI
2 GAUZE CLEAN-WRAP BDGE N/S 2"
1 GAUZE CLEAN-WRAP BDGE N/S 3"
1 ABD COMBINE PAD 5" X 9"
1 CPR FILTERSHIELD 77-100
1 BAGGED COMP MISC
1 1 OZ, BUFF EYEWASH
1 SCISSOR BDGE 4" RED PLS HDL
LBL STOCK 6-3/8"X4"
LBL STOCK 4"X2-7/8"
1 LBL STOCK 3"x1-7/8"
2 PR LRG NITRILE GLVES ZIP BAG
2 1" X 3" PLASTIC BANDS 16/BAG
2 TAPE ADHESIVE 1/2 X 2.5 125133
1 ADH BNDG PLASTIC EX-LG 4"X 2"
1 KIT, PP 16 UNIT FA
1 LBL CONTENTS ANSI Z308.1-2009 REV B
1 TRI BNDG NON WOVEN 40"X40"X56"
1 COLD PACK UNIT 4"X6" BULK
4 GAUZE PADS 2"X2" 12PLY
1 EYE PADS STD OVAL STERILE
3 WOVEN FINGERTIP BANDAGE 2"
2 WOVEN KNUCKLE BANDAGE

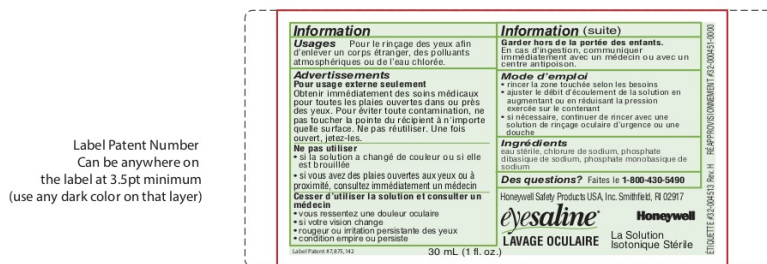
Eye Wash Package label

#32-004513 Rev. H



Top Panel
3/32" from die edges

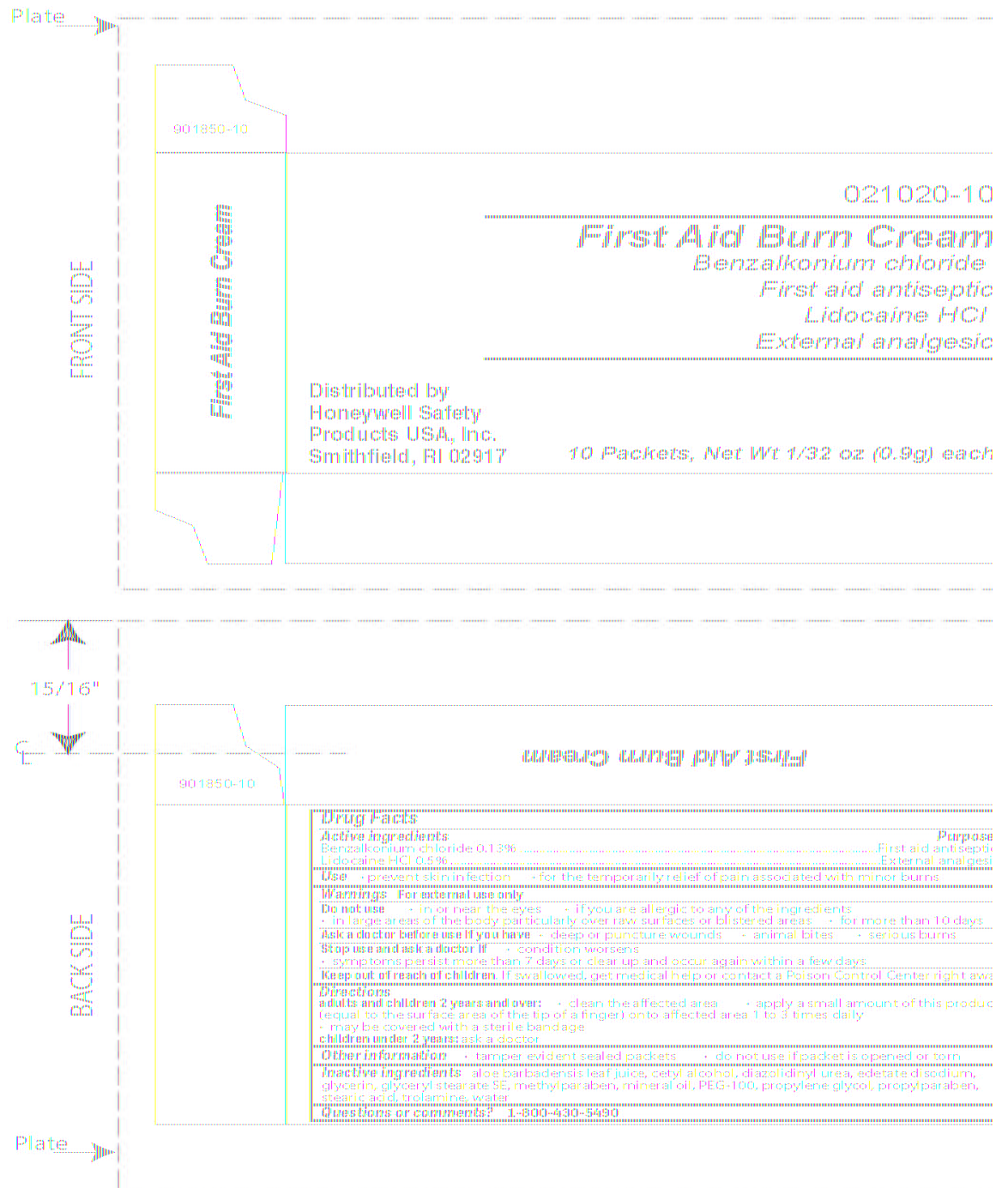
Back
1/8" from die edges



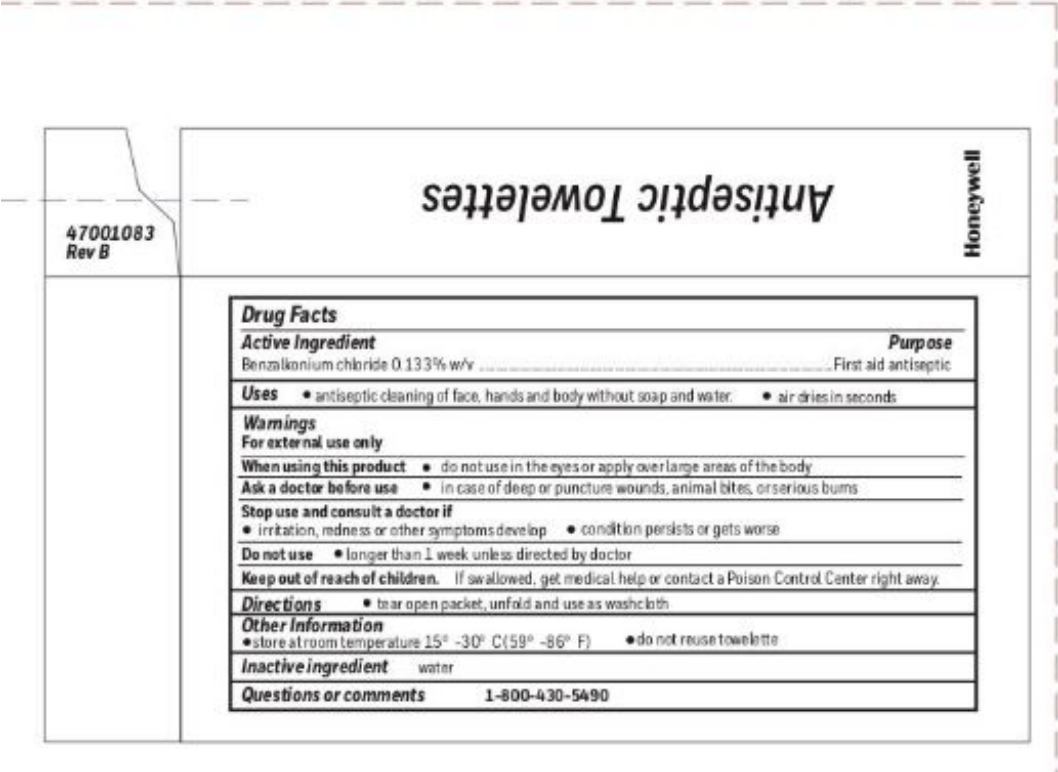
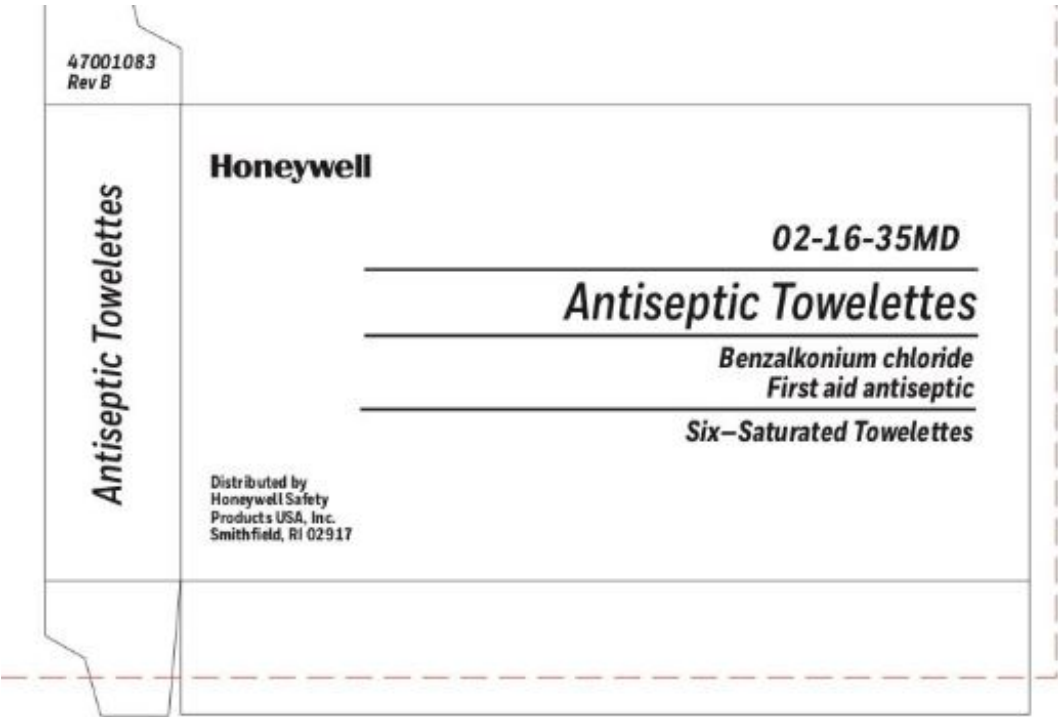
Base
3/32" from die edges

Printable Text Area

First Aid Burn Cream
Principal Display Panel



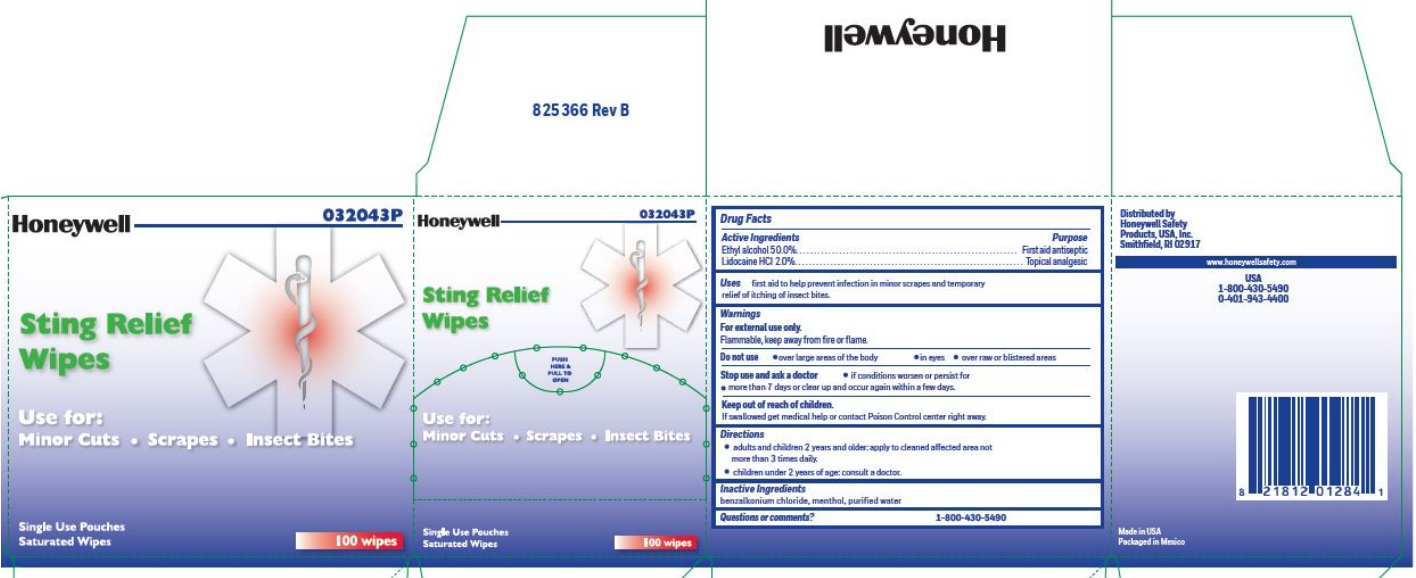
Principal Display Panel



Aypanal
Principal Display Panel



Sting Relief Principal Display Panel



Neomycin Antibiotic Ointment Principal Display Panel

822568-25

**Neomycin First
aid antibiotic**

020126-25

Neomycin
First aid antibiotic
Neomycin sulfate
First aid antibiotic

Distributed by:
Honeywell Safety Products
USA, Inc. Smithfield,
RI 02917

25 Packets, Net Wt 1/32 oz (0.9 g) each

Neomycin First aid antibiotic

822568-25

Drug Facts

Active ingredient (in each gram)	Purpose
Neomycin sulfate (equivalent to 3.5 mg neomycin)	First aid antibiotic

Use first aid to help prevent infection in minor • cuts • scrapes • burns

Warnings

For external use only

Allergy alert • do not use if you are allergic to any of the ingredients

Do not use • in the eyes • over large areas of the body

Ask a doctor before use if you have • deep or puncture wounds • animal bites • serious burns

Stop use and ask a doctor if • conditions persists or gets worse • rash or other allergic reaction develops
• you need to use longer than one week

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions • clean the affected area

• apply a small amount of product (equal to the surface area of the tip of a finger) on the area 1 to 3 times daily
• may be covered with a sterile bandage

Other information store at 15° to 25°C (59° to 77°F)

Inactive ingredient petrolatum

Questions or comments? 1-800-430-5490

4132 Kit Label
SF00003655

Cover Size: 9.75"w x 6.5"h

TYPE OF KIT: Plastic 16 Unit Kit

SCREEN NUMBER: SF00003655-5194

LOGO SIZE: 8.88" x 5.50"

PRINTED IN: PMS Red 485 and PMS 349 Green



FIRST AID

FOR REFILL CALL – (888) 259-7788

Distributed by Honeywell Safety Products USA, Inc. Smithfield, RI 02917

If distributor is signing for end user approval of logo, distributor assumes responsibility.

This drawing has been reviewed and approved for the following: kit size, ink color(s), kit part number and logo being printed on the kit

☐ Needs changes

☐ Approved

(A digital signature or response via email with attached approval are both accepted as an approval to this form. Please indicate any changes needed)

Name

Signature

Date

FRM-03-052 Rev. *

(8/18/10)

1/1

4132 FIRST AID KIT

4132 first aid kit kit

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0498-4132
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Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0498-4132-01	1 in 1 KIT; Type 0: Not a Combination Product	10/18/2018	

Quantity of Parts

Part #	Package Quantity	Total Product Quantity
Part 1	1 BOTTLE	30 mL
Part 2	3 PACKET	6
Part 3	6 POUCH	2.4 mL
Part 4	10 PACKET	9 g
Part 5	10 PACKET	9 g
Part 6	10 PACKET	14 mL

Part 1 of 6

EYESALINE EMERGENCY EYEWASH

purified water liquid

Product Information

Item Code (Source)	NDC:0498-0100
Route of Administration	OPHTHALMIC

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
WATER (UNII: 059QF0KO0R) (WATER - UNII:059QF0KO0R)	WATER	98.6 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
SODIUM PHOSPHATE, DIBASIC (UNII: GR686LBA74)	
SODIUM PHOSPHATE, MONOBASIC, MONOHYDRATE (UNII: 593YOG76RN)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0498-0100-01	30 mL in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part349	12/18/2018	

Part 2 of 6

AYPANAL NON-ASPIRIN

acetaminophen tablet

Product Information

Item Code (Source)	NDC:0498-2001
Route of Administration	ORAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	325 mg

Inactive Ingredients

Ingredient Name	Strength
STEARIC ACID (UNII: 4ELV7Z65AP)	
SODIUM STARCH GLYCOLATE TYPE A CORN (UNII: AG9B65PV6B)	
STARCH, CORN (UNII: O8232NY3SJ)	
POVIDONE (UNII: FZ989GH94E)	

Product Characteristics

Color	white	Score	2 pieces
Shape	ROUND	Size	10mm
Flavor		Imprint Code	circle;U
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0498-2001-01	2 in 1 PACKET; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part343	04/10/2012	

Part 3 of 6

STING RELIEF PAD

ethyl alcohol, lidocaine swab

Product Information

Item Code (Source)	NDC:0498-0733
Route of Administration	TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LIDOCAINE HYDROCHLORIDE (UNII: V13007Z41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE ANHYDROUS	20 mg in 1 mL
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	0.5 mL in 1 mL

Inactive Ingredients

Ingredient Name	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
MENTHOL (UNII: L7T10EIP3A)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1		0.4 mL in 1 POUCH; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	12/23/2017	

Part 4 of 6

FIRST AID BURN

benzalkonium chloride, lidocaine hydrochloride cream

Product Information

Item Code (Source)	NDC:0498-0903
Route of Administration	TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.13 g in 100 g
LIDOCAINE HYDROCHLORIDE (UNII: V13007Z41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE	0.5 g in 100 g

Inactive Ingredients	
Ingredient Name	Strength
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
DIAZOLIDINYL UREA (UNII: H5RIZ3MPW4)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
WATER (UNII: 059QF0KO0R)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	
PEG-100 STEARATE (UNII: YD01N1999R)	
LIGHT MINERAL OIL (UNII: N6K5787QVP)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
TROLAMINE (UNII: 9O3K93S3TK)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1		0.9 g in 1 PACKET; Type 0: Not a Combination Product		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	12/20/2017	

Part 5 of 6
NEOMYCIN antibiotic ointment

Product Information	
Item Code (Source)	NDC:0498-0730
Route of Administration	TOPICAL

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
NEOMYCIN SULFATE (UNII: 057Y626693) (NEOMYCIN - UNII:I16QD7X297)	NEOMYCIN SULFATE	3.5 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
PETROLATUM (UNII: 4T6H12BN9U)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0498-0730-01	0.9 g in 1 PACKET; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part333B	03/31/2010	

Part 6 of 6

ANTISEPTIC TOWELETTE

benzalkonium chloride liquid

Product Information

Item Code (Source)	NDC:0498-0501
Route of Administration	TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	1.3 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1		1.4 mL in 1 PACKET; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	12/21/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		10/18/2018	

Labeler - Honeywell Safety Products USA, INC (079287321)

Establishment

Name	Address	ID/FEI	Business Operations
Honeywell Safety Products USA, INC		079287321	pack(0498-4132)

Establishment

Name	Address	ID/FEI	Business Operations
Ultra Seal Corporation		085752004	manufacture(0498-2001)

Establishment

Name	Address	ID/FEI	Business Operations
Water-Jel Technologies		155522589	manufacture(0498-0903, 0498-0730)

Establishment

Name	Address	ID/FEI	Business Operations
Honeywell Safety Products USA, Inc		167518617	manufacture(0498-0100)

Establishment

Name	Address	ID/FEI	Business Operations
Changzhou Maokang Medical		421317073	manufacture(0498-0501)

Establishment

Name	Address	ID/FEI	Business Operations
Safetec of America Inc		874965262	manufacture(0498-0733)

Revised: 4/2019

Honeywell Safety Products USA, INC